Estimation of rutin and quercetin Terminalia chebula by HPLC

A Kumar, K Lakshman, K Jayaveera, K Satish, S Tripathi

Citation

A Kumar, K Lakshman, K Jayaveera, K Satish, S Tripathi. *Estimation of rutin and quercetin Terminalia chebula by HPLC*. The Internet Journal of Aesthetic and Antiaging Medicine. 2008 Volume 2 Number 1.

Abstract

Tannins and Flavonoids present in the Terminalia chebula, Flavonoids like Rutin and quercetin possess many biochemical effects like inhibition of enzymes, regulatory role on different hormones and pharmacological activities like antimicrobial, antioxidant, anticancer, antihepatotoxic, protection of cardio vascular system. An HPLC method was developed for the estimation of rutin and quercetin from methanol methanolic extract of Terminalia chebula

INTRODUCTION

Terminalia chebula is one of the ingredients present in the many ayurvedic and other traditional medicine system. Terminalia chebula is traditionally used in formulation for anti-diabetic, anti-inflammatory, laxative, antibacterial, antifungal, cardiotonic, diuretic, hyperlipidemic activity, jaundice (Anonymous. 1999, Kirtikar KR and Basu BD, 1987, Inamdar et al., 1959, Sabu, M.C. 2002 Miglani, et al., 1971, Khanna, et al., 1993). Flavonoids are a group of polyphenolic compounds, which are widely distributed through out the plant kingdom. To date about 300 varieties of flavonoids are known (Anonymous, 1996). Many have low toxicity in mammals and some of them are widely used in medicine for maintenance of capillary integrity (Kuhnau, J. 1976). Rutin, 5,7,3', 4',tetrahydroxy flavonol -3rhamnoglucoside and quercetin 5,7,3',4',- tetrahydroxy flavonol are exhibits anti-inflammatory, antihepatotoxic (Cesarone, 1992), antiulcer (Clack et al., 1950), antiallergic, antiviral actions and some of them provides protection against cardiovascular mortality (Colergie Smith et al., 1980, Hertog et al., 1993). Both possess antioxidant activity and reduce low density lipoproteins (LDL) oxidation (Dewhalley et al., 1990), quercetin in combination with other flavonoids, are inhibiting a number of enzymes like bradykinin (Bamard et al., (1993), tyrosine kinase (Hur, et al., (1994), and 5'-nucleotidase activity (Beladi, et al., 1987). Rutin and quercetin have shown regulatory activity of hormones like affect the transport, metabolism and action of thyroid hormones. High performance layer chromatography (HPLC) (Harbone, J.B. 1984) method is the suitable method

for estimation of chemical constituents present in plant materials. Hence Terminalia chebula contains rutin and quercetin are important active constituents and is estimated by HPLC method.

MATERIALS AND METHODS INSTRUMENTATION

The Shimadzu class LC-10AT HPLC, Hichrom C18 and a Rheodyne 7725i injector fitted with a 20 ll loop, column oven, and a photodiode array detector. The output signal was monitored and processed using chromquest version3.0 software on Pentium computer (Hewlett Packard).

SOLVENT AND CHEMICALS

Rutin and quercetin obtained from natural remedies (Bangalore) chromatographic grade methanol, formic acid and acetonitrile (AR), were obtained from Merck (Mumbai, India).

PLANT MATERIAL

Terminalia chebula fruits were extracted with distilled water by Soxhlet apparatus. The pooled aqueous extract was evaporated under vacuum to dryness, yielding was noted. Aqueous extract was subjected for estimation of rutin and quercetin.

PREPARATION OF STANDARD AND SAMPLE SOLUTIONS

Rutin and quercetin 10 mg were accurately weighed into a 10 mL volumetric flask, dissolved in 5 mL methanol and the solution was made up to 10 mL with the same solvent (1

mg/mL). T. Chebula fruit extract was accurately weighed (10 mg) into a 10 mL of volumetric flask and dissolved in methanol the solution was filtered through Whatman filter paper No. 42 and the filtrate was made up to the mark with methanol.

{image:1}

RESULTS AND DISCUSSION

The retention time of standards rutin and quercetin were found to be 4.072 and 19.104 (Graphs1). The retention time of rutin and quercetin in Terminalia chebula were found to be 4.802 and 19.040 (Graphs2), which are matching with standard R_t values respectively. Then the amount of rutin and quercetin in Terminalia chebula was found to be 59.52 and 9.06 % w/v respectively graphs

{image:2}

{image:3}

ACKNOWLEDGEMENT

The authors are thankful to K.V.Naveen Kiran, Chairman, Sri K.V.College of Pharmacy, Chickballapur, for providing required facilities.

References

- 1. Anonymous. (1999). Indian Herbal Pharmacopoeia. Regional Research Laboratory, Jammutwai and Indian Drug Manufacturers Association Mumbai, 2, (1999) pp.51. 2. Kirtikar, K.R., and Basu, B.D., (1987). Indian Materia Medica, International book distributors, Dehra Dun, India, 3, pp. 333-335.
- 3. Inamdar, M.C., Khorana, M.L., Rao M.R.R., (1959). Antibacterial and antifungal activity of Terminalia chebula Retz. Indian Journal of Pharmacy; 21(12); 333-335.
- 4. Sabu, M.C. (2002). Ramadasan Kuttan. Anti-diabetic

- activity of medicinal plants and its relationship with their antioxidant property, Journal of Ethanopharmacology. 81 (2); 155-160.
- 5. Miglani, B.D., Sen, P., and Sanyal, R.K. (1971). Purgative action of an oil obtained from Terminalia chebula, Indian Journal of Medical Research, , 59 (2); 281-283.
- 6. Khanna, A.K., Chander, R., Kapoor, N.K., Singh, C., Srivastava. A.K. (1993). Hypolipidemic activity of Terminalia chebula in rats, Fitoterapia. 64 (4); 351-356.
- 7. Anonymous, (1996). Indian Pharmacopoeia, Vol II. Ministry of Health & Family Welfare. Govt. of India, Controller of Publications, New Delhi, 53 - 54.
- 8. Kuhnau, J. (1976). The flavonoids: A class of semiessential food components: their role in human nutrition. World Res Nut Diet, 24; 17-91.
- 9. Cesarone, M.R., Laurora, G., Ricci, A., Belcaco, G., Pomante, P. (1992). Acute effects of hydroxyethylrutosides on capillary filtration in normal volunteers, patients with various hypotensions and in patients with diabetic micro angiopathy. J Vas Disease; 21; 76-80.
- 10. Clack, W., Heller, W., Michel, C., Saran, M., (1950). Effect of flavonoid substances on histamine toxicity, anaphylactic shock and histamine-enhanced capacity to dye J Allergy, 21; 133-147.
- 11. Colergie Smith, P.O., Thomas, P., Scurry, J.H., Dormandy, J.A., (1980). Causes of various ulceration, a New Hypothesis. Br Med J, 296; 1726-7.
- 12. Hertog, M.G.L., Hollman, P.C.H., Katan, Klohout, D. (1993). Intake of potentially anticarcinogenic flavonoids and their determinants in adults in the Netherlands. Nutr Cancer;
- 13. De-whalley, C., Rankin, S.M., Houct, J.R.S., Jessup, W., Leake, D.S. (1990). Flavonoids inhibit the oxidative modification of low-density lipoproteins by macrophages. Biochem Pharmacol, 39; 1743-50.
- 14. Bamard, D.L., Smee, D.F., Huffman, J.H., Meyerson, C.R., Sidwell, R.W. (1993). Review of Quercetin and related
- bioflavonoids. Chemotherapy, 39; 203-11. 15. Hur, C.Q., Chen, K., Shi, Q., Kikushkie, R.E., Cheng, Y.C., Lee, K.H. (1994). Apoptosis of HIV infected cell following treatment with Shosoikorso. J Nat Pro, 57; 42-50. 16. Beladi, I., Musci, R., Pusztai, M., Bakay, I., Rosztoczy, M., Gabor. (1987). Bioactivity of flavonoids. Stud Org Chem, 57; 42-50.
- 17. Harbone, J.B. (1984). Phytochemical Methods, Chapman and Hall, 2nd edition, New York, 1-31.

Author Information

Ashok Kumar, B.S, M.Pharma, Ph.D., Scholar

Department of Pharmacognosy, Sri K.V.College of Pharmacy

K. Lakshman, M.Pharma, Ph.D.

Department of Pharmacognosy, PES College of Pharmacy

K.N. Jayaveera

Department of Chemistry, JNTU College of Engineering

K.V. Satish, M.Pharma, Ph.D., Scholar

Department of Pharmaceutics, Sri K.V.College of Pharmacy

Sachidanand Mani Tripathi, M.Pharma

Department of Pharmacognosy, Sri K.V.College of Pharmacy